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WHAT IS CLAIMED IS:

1. An isolated and purified polynucleotide comprising a nucleic acid sequence encoding a WWOX polypeptide.

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The polynucleotide of claim 1, comprising a nucleic acid sequence encoding SEQ 2. ID NO:2.

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The polynucleotide of claim 1, comprising a nucleic acid sequence encoding SEQ ID NO:31.

4. The polynucleotide of claim 1, comprising a nucleic acid sequence encoding SEO ID NO:33.

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The polynucleotide of claim 2, comprising SEQ ID NO:30. VV

The polynucleotide of claim 2, comprising SEO ID NO:1.

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The polynucleotide of claim 2, comprising SEQ ID NO:32. 7.

8. The polynucleotide of claim 1, comprising a nucleic acid sequence encoding at least 50 contiguous amino acid residues of SEQ ID NO:2.

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9. The polynucleotide of claim 8, comprising a nucleic acid sequence encoding at least 150 contiguous amino acid residues of SEQ ID NO:2.

10. The polynucleotide of claim 1, comprising at least 1.5 contiguous kilobases of SEQ ID NO:1.

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An expression vector comprising a nucleic acid sequence encoding a WWOX 11. polypeptide.

- 12. The expression vector of claim 11, wherein the nucleic acid sequence encodes SEQ ID NO:2.
- 5 13. The expression vector of claim 11, wherein the nucleic acid sequence encodes SEQ ID NO:31.
 - 14. The expression vector of claim 11, wherein the nucleic acid sequence encodes SEQ ID NO:33.

- 15. The expression vector of claim 11, wherein the nucleic acid sequence comprises SEQ ID NO:1.
- 16. The expression vector of claim 11, wherein the nucleic acid sequence comprises15 SEQ ID NO:30.
 - 17. The expression vector of claim 11, wherein the nucleic acid sequence comprises SEQ ID NO:32.
- 20 18. The expression vector of claim 11, wherein the nucleic acid sequence comprises at least 1.5 contiguous kilobases of SEQ ID NO:1.
 - 19. The expression vector of claim 18, wherein the nucleic acid sequence encodes at least 50 contiguous amino acids of SEQ ID NO:2.

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- 20. The expression vector of claim 11, wherein the nucleic acid sequence further comprises a promoter operably linked to the WWOX-encoding nucleic acid sequence.
- 21. The expression vector of claim 20, wherein the promoter is heterologous.

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- 22. The expression vector of claim 20, wherein the promoter is a constitutive promoter, a tissue-specific promoter, an inducible promoter, or a noninducible promoter.
- 23. The expression vector of claim 11, wherein the expression vector is a viral vector.

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- 24. The expression vector of claim 23, wherein the viral vector is a vaccinia virus, adenovirus, herpesvirus, retrovirus, cytomegalovirus, or adeno-associated virus.
- 25. A recombinant host cell comprising a nucleic acid sequence encoding a WWOX 10 polypeptide.
 - 26. The recombinant host cell of claim 25, wherein the polypeptide comprises SEO ID NO:2.

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27. The recombinant host cell of claim 25, wherein the polypeptide comprises SEQ ID NO:31.

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28. The recombinant host cell of claim 25, wherein the polypeptide comprises SEQ ID NO:33.

The recombinant host cell of claim 25, wherein the nucleic acid sequence comprises

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- SEQ ID NO:1.
- 30. The recombinant host cell of claim 25, wherein the nucleic acid sequence comprises 25 SEQ ID NO:30.
 - 31. The recombinant host cell of claim 25, wherein the nucleic acid sequence comprises SEQ ID NO:32.
- 30 32. A method of preparing recombinant WWOX comprising:

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- (a) transfecting a cell with a polynucleotide comprising a nucleic acid sequence encoding a WWOX polypeptide to produce a transformed host cell; and
- (b) maintaining the transformed host cell under biological conditions sufficient for expression of the WWOX polypeptide in the host cell.
- 33. The method of claim 32, wherein the nucleic acid sequence encodes SEQ ID NO:2.
- 10 34. The method of claim 32, wherein the nucleic acid sequence encodes SEQ ID NO:31.
 - 35. The method of claim 32, wherein the nucleic acid sequence encodes SEQ ID NO:33.
 - 36. The method of claim 32, wherein the nucleic acid sequence comprises SEQ ID NO:1.
 - 37. The method of claim 32, wherein the nucleic acid sequence comprises SEQ ID NO:30.
 - 38. The method of claim 32, wherein the nucleic acid sequence comprises SEQ ID NO:32.
- 25 39. The method of claim 32, wherein the polynucleotide is comprised in a vector.
 - 40. A method of treating a pre-cancer or cancer cell comprising providing to the cell an amount of a WWOX polypeptide effective to induce apoptosis in the cell.

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- 41. The method of claim 40, wherein the WWOX polypeptide is provided to the cell by administering an expression vector comprising a polynucleotide encoding a WWOX polypeptide under the transcriptional control of a promoter.
- The method of claim 40, wherein the cell is a bladder, blood, bone, bone marrow, brain, breast, central nervous system, colon, esophagus, gastrointestine, head, kidney, liver, lung, nasopharynx, neck, ovary, prostate, skin, stomach, or uterus cell.
 - 43. The method of claim 40, wherein the expression vector comprises a viral vector.
 - 44. A method of treating a subject having a hyperproliferative condition comprising contacting a cell within the subject with an expression vector comprising a polynucleotide encoding an WWOX polypeptide under the transcriptional control of a promoter, wherein expression of the WWOX polypeptide confers a therapeutic benefit on the subject.
 - 45. The method of claim 44, wherein the cell is a cancer or pre-cancer cell.
 - 46. The method of claim 44, wherein the cell is involved with restenosis, primary psoriasis, angiogenesis, rheumatoid arthritis, inflammatory bowel disease, psoriasis, eczema, secondary cataracts, or bronchial dysplasia.
 - 47. The method of claim 45, wherein the cancer or pre-cancer cell is selected from a group consisting of a bladder, blood, bone, bone marrow, brain, breast, colon, esophagus, gastrointestine, head, kidney, liver, lung, nasopharynx, neck, ovary, prostate, skin, stomach, and uterus cell.
 - 48. The method of claim 45, wherein the cancer or pre-cancer cell is derived from or is part of a solid tumor.
 - 49. The method of claim 44, wherein the contacting occurs in vitro.

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- 50. The method of claim 44, wherein the contacting occurs in vivo.
- 51. The method of claim 44, wherein the expression vector is delivered endoscopically, intravenously, intralesionally, percutaneously, or subcutaneously.
 - 52. The method of claim 48, wherein the expression vector is delivered by direct injection into the tumor.
- 10 53. The method of claim 44, wherein the expression vector comprises a viral vector.
 - 54. The method of claim 53, wherein the viral vector a vaccinia virus, adenovirus, herpesvirus, retrovirus, cytomegalovirus, or adeno-associated virus.
- 15 55. The method of claim 44, wherein the contacting is performed at least twice.
 - 56. The method of claim 55, wherein the second contacting follows the first by a period of about one day to one year.
- 57. The method of claim 48, further comprising contacting the tumor with an anticancer therapy.
 - 58. The method of claim 57, wherein the anticancer treatment is chemotherapy, immunotherapy, surgery, radiotherapy, gene therapy with a second therapeutic polynucleotide other than a polynucleotide encoding the WWOX polypeptide, or other biotherapy.
 - 59. The method of claim 57, wherein the expression vector is contacted with the tumor prior to, at the same time as, or after contacting with the anticancer treatment.

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- 60. The method of claim 44, wherein the endogenous WWOX polypeptide of the cancer cell is mutated.
- 61. A method for detecting the susceptibility of an individual to a certain cancer comprising;
 - (i) obtaining DNA from an individual;
 - (ii) obtaining probes specific to WWOX; and
 - (ii) identifying a change in the WWOX gene and/or gene products.
- 10 62. The method of claim 61, wherein the identifying comprises amplification.
 - 63. The method of claim 62, wherein the probes encode nucleic acid primers.
 - 64. The method of claim 61, wherein the change is a mutation of WWOX.
 - 65. The method of claim 61, wherein the change is a increase in the amount of a WWOX gene product.
 - 66. The method of claim 61, wherein the change is a decrease in the amount of a WWOX gene product.
 - 67. The method of claim 61, wherein the DNA is genomic DNA.
 - 68. The method of claim 67, wherein the genomic DNA is chromosomal DNA.
 - 69. The method of claim 68, wherein the identifying comprises fluorescent *in situ* hybridization.
- 70. The method of claim 69, wherein the probes encode nucleic acids spanning the WWOX chromosomal locus.

- 71. The method of claim 70, wherein the probes further comprise a fluorescent detection moiety.
- 72. The method of claim 61, wherein the cancer is multiple myeloma.
- 73. The method of claim 61, wherein the cancer is breast cancer.